

Amendments to the Specification:

Please replace the paragraph inserted before the first line of the specification as requested on December 6, 2001 with the following amended paragraph:

This application is a Divisional patent application [~~under 37 C.F.R. § 1.53(b)~~] of [Continuation] U.S. Application No. 09/506,729, now U.S. Patent 6,365,352, filed on February 18, 2000 [~~(pending allowance)~~], which [~~claims priority to~~] is a continuation of PCT International Application No. PCT/US98/17284, filed August 21, 1998, which claims [~~priority to~~] the benefit of U.S. Provisional Application No. 60/056,844, filed on August 22, 1997, all of which are herein incorporated by reference in their entirety.

Please insert the following paragraph before “**Technical Field**” on page 1, line 5 of the specification:

This application is related to application serial No. 08/510,032, serial No. 60/056,844 and application serial No. 08/688,514, all of which are herein incorporated by reference in their entirety. All published articles, patents and other publications cited throughout this application are herein incorporated by reference in their entirety.

Please replace the paragraph beginning on page 9, line 7, with the following amended paragraph:

Figs. 3A and B Figure 3 is an autoradiogram of the expression profile generated from cDNAs made with RNA isolated from neutrophils exposed to avirulent *E. coli* and virulent and avirulent *Y. pestis*. All possible 12 anchoring oligo d(T)n1, n2 were used to generate a complete expression profile for the enzyme *Bgl*II.

Please replace the paragraph beginning on page 13, line 23, with the following amended paragraph:

The oligonucleotide primer that primes first strand DNA synthesis comprises a 5' sequence incapable of hybridizing to a polyA tail of the mRNAs, and a 3' sequence that hybridizes to a portion of the polyA tail of the mRNAs and at least one non-polyA nucleotide immediately upstream of the polyA tail. The 5' sequence is preferably a sufficient length that can serve as a primer for amplification. The 5' sequence also preferably has an average G+C content and does not contain large palindromic sequence; some palindromes, such as a recognition sequence for a restriction enzyme, may be acceptable. Examples of suitable 5' sequences are

CTCTCAAGGATC[:]TACCGCT (SEQ ID [No.] NO: 1),

CAGGGTAGACGACGCTACGC (SEQ ID [No.] NO: 2), and

TAATACCGCGCCACATAGCA (SEQ ID [No.] NO: 3).

Please replace the paragraph beginning on page 15, line 14, with the following amended paragraph:

The adapters for use in the present invention are designed such that the two strands are only partially complementary and only one of the nucleic acid strands that the adapter is ligated to can be amplified. Thus, the adapter is partially double-stranded (*i.e.*, comprising two partially hybridized nucleic acid strands), wherein portions of the two strands are non-complementary to each other and portions of the two strands are complementary to each other. Conceptually, the adapter is “Y-shaped” or “bubble-shaped.” When the 5’ region is non-paired, the 3’ end of other strand cannot be extended by a polymerase to make a complementary copy. The ligated adapter can also be blocked at the 3’ end to eliminate extension during subsequent amplifications. Blocking groups include dideoxynucleotides or any other agent capable of blocking the 3’-OH. In this type of adapter (“Y-shaped”), the non-complementary portion of the upper strand of the adapters is preferably a length that can serve as a primer for amplification. As noted above, the non-complementary portion of the lower strand need only be one base, however, a longer sequence is preferable (*e.g.*, 3 to 20 bases; 3 to 15 bases; 5 to 15 bases; or 14 to 24 bases). The complementary portion of the adapter should be long enough to form a duplex under conditions of li[ti]gation.

Please replace the paragraph beginning on page 23, line 8, with the following amended paragraph:

Synthesis of cDNA was performed as previously described by Prashar *et al.* in WO 97/05286 and in Prashar *et al.* (1996) *Proc. Natl. Acad. Sci. USA* 93:659-663. Briefly, cDNA was synthesized according to the protocol described in the GIBCO/BRL kit for cDNA synthesis. The reaction mixture for first-strand synthesis included 6 µg of total RNA, and 200 ng of a mixture of 1-base anchored oligo(dT) primers with all three possible anchored bases (ACGTAATACGACTCACTATAGGGCGAATTGGGTCGACTTTTTTTTTTTn1 wherein n1=A/C or G, SEQ ID NO: 4) along with other components for first-strand synthesis reaction except reverse transcriptase. This mixture was incubated at 65°C for 5m, chilled on ice and the process repeated. Alternatively, the reaction mixture may include 10µg of total RNA, and 2 pmol of 1 of the 2-base anchored oligo(dT) primers a heel such as RP5.0 (CTCTCAAGGATCTTACCGCTT₁₈AT, SEQ ID NO: 5), or RP6.0 (TAATACCGCGGCCACATAGCAT₁₈CG, SEQ ID NO: 6), or RP9.2 (CAGGGTAGACGACGCTACGCT₁₈GA, SEQ ID NO: 7) along with other components for first-strand synthesis reaction except reverse transcriptase. This mixture was then layered with mineral oil and incubated at 65°C for 7 min followed by 50°C for another 7 min. At this stage, 2µl of Superscript reverse transcriptase (200 units/µl; GIBCO/BRL) was added quickly and mixed, and the reaction continued for 1 hr at 45-50°C. Second-strand synthesis was performed at 16°C for 2 hr. At the end of the reaction, the cDNAs were precipitated with ethanol and the yield of cDNA was calculated. In our experiments, ≈200 ng of cDNA was obtained from 10µg of total RNA.

Please replace the paragraph beginning on page 24, line 1, with the following amended paragraph:

The adapter oligonucleotide sequences were A1 (TAGCGTCCGGCGCAGCGACGGCCAG, SEQ ID NO: 8) and A2 (GATCCTGGCCGTGGCTGTCTGTCGGCGC, SEQ ID NO: 9). One microgram of oligonucleotide A2 was first phosphorylated at the 5' end using T4 polynucleotide kinase (PNK). After phosphorylation, PNK was heated denatured, and 1µg of the oligonucleotide A1 was

added along with 10× annealing buffer (1 M NaCl/100 mM Tris-HCl, pH8.0/10 mM EDTA, pH8.0) in a final vol of 20 µl. This mixture was then heated at 65°C for 10 min followed by slow cooling to room temperature for 30 min, resulting in formation of the Y adapter at a final concentration of 100 ng/µl. About 20 ng of the cDNA was digested with 4 units of *Bgl* II in a final vol of 10 µl for 30 min at 37°C. Two microliters (\approx 4 ng of digested cDNA) of this reaction mixture was then used for ligation to 100 ng (\approx 50-fold) of the Y-shaped adapter in a final vol of 5µl for 16 hr at 15°C. After ligation, the reaction mixture was diluted with water to a final vol of 80 µl (adapter ligated cDNA concentration, \approx 50 pg/µl) and heated at 65°C for 10 min to denature T4 DNA ligase, and 2-µl aliquots (with \approx 100 pg of cDNA) were used for PCR.

Please replace the paragraph beginning on page 24, line 17, with the following amended paragraph:

The following sets of primers were used for PCR amplification of the adapter ligated 3' -end cDNAs:

TGAAGCCGAGACGTCGGTCG(T)₁₈ n1, n2 (wherein n1, n2 = AA, AC, AG AT CA CC CG CT GA GC GG and GT; SEQ ID NO: 10) as the 3' primer with A1 as the 5' primer or alternatively

RP 5.0, RP 6.0, or RP 9.2 used as 3' primers with primer A1.1 serving as the 5' primer. To detect the PCR products on the display gel, 24 pmol of oligonucleotide A1 or A1.1 was 5' -end-labeled using 15 µl of [γ -³² P]ATP (Amersham; 3000 Ci/mmol) and PNK in a final volume of 20 µl for 30 min at 37°C. After heat denaturing PNK at 65°C for 20 min, the labeled oligonucleotide was diluted to a final concentration of 2 µM in 80 µl with unlabeled oligonucleotide A1.1. The PCR mixture (20µl) consisted of 2 µl (\approx 100 pg) of the template, 2µl of 10× PCR buffer (100 mM Tris-HCl, pH 8.3/500 mM KCl), 2 µl of 15 mM MgCl₂ to yield 1.5 mM final Mg²⁺ concentration optimum in the reaction mixture, 200 µM dNTPs, 200 nM each 5' and 3' PCR primers, and 1 unit of AmpliTaq Gold. Primers and dNTPs were added after preheating the reaction mixture containing the rest of the components at 85°C. This “hot start” PCR was done to avoid artefactual artifactual amplification arising out of arbitrary annealing of PCR primers at lower temperature during transition from room temperature to 94°C in the first PCR cycle. PCR consisted of 5 cycles of 94°C for 30 sec, 55°C for 2 min, and 72°C for 60 sec followed by 25

cycles of 94°C for 30 sec, 60°C for 2 min, and 72°C for 60 sec. A higher number of cycles resulted in smeary gel patterns. PCR products (2.5µl) were analyzed on 6% polyacrylamide sequencing gel. For double or multiple digestion following adapter ligation, 13.2 µl of the ligated cDNA sample was digested with a secondary restriction enzyme(s) in a final vol of 20 µl. From this solution, 3µl was used as template for PCR. This template vol of 3 µl carried ≈ 100 pg of the cDNA and 10 mM MgCl₂ (from the 10× enzyme buffer), which diluted to the optimum of 1.5 mM in the final PCR vol of 20 µl. Since Mg²⁺ comes from the restriction enzyme buffer, it was not included in the reaction mixture when amplifying secondarily cut cDNA. Bands were extracted from the display gels as described by Liang *et al.* (1995 *Curr. Opin. Immunol.* 7:274-280), reamplified using the 5' and 3' primers, and subcloned into pCR-Script with high efficiency using the PCR-Script cloning kit from Stratagene. Plasmids were sequenced by cycle sequencing on an ABI automated sequencer.

Please replace Table 2, beginning on page 37 with the following Table 2.

TABLE 2

Cln	Sequence
846	1 TCTCAGTGAG CTGAGATCAC ACCACTGCAC TCCAACTGGG CGACAGAGCA 51 AG <u>(SEQ ID NO: 11)</u>
854	1 CACTTCCCC AAATTCTTT GCCATAGTTC ACTCTCTACT GATAAGGCCA 51 C. <u>(SEQ ID NO: 12)</u>

855	1 GGGAAAGTGG TGGGGTGGTG AGGGTCAATG TGCAGAAAAT CGATGTAACT 51 TGTAATACAG TTGAGTCAAC TGTGTGTTCA CAACAACCTCT GAGAGTTAAC 101 ACCATTCTA C (<u>SEQ ID NO: 13</u>)
856	1 ATCTAAATAT TTTCATACC GAGTTATTAA GGAGTCAGTA GTCTGTGCTA 51 CAATGCTGCA AAAAGCATCA CGTGGAAAGAA TGGGAACATAT GCGTACTTTA 101 TGAAGTGATG TATAACACAA TGAACCTCTGT TTTACAACTA CAGTGCTGCA 151 TTCAATTATC TTCCAT (<u>SEQ ID NO: 14</u>)
859	1 AAGCTCTGTA TACAAAAGTT ATTTATTTAG ATGTTCGAGG CATGTCTCTC 51 CTCACCTGTA AACTAACTGT TTTATAACAG CTTGTATCAC ATGTGTGAAG 101 TTAATGAATG TAATACTCCA ACAAGCCATT CATCAGATTG GCCAACAGCT 151 AGGATACAGT TAAATAATGG CGACCAGGTT GACAAGTCAT AATTGCGGTT 201 TGGGGGACCG TAGTTGCACC TCACCTAGAC CAACGTACGC ATGGCACTCG 251 ACCCAGGCGA ACAAAATTAA T (<u>SEQ ID NO: 15</u>)

863	<p>1 TTTCTCAAGA AGAGATAAGA ATGAAAAGTC ATAGAACACA TCATGGAGGA</p> <p>51 CCTGGACACA AATGCAGACA AGCAGCTGAG CTTCGAGGAG TTCATCATGC</p> <p>101 TGATGGCGAG GCTAACCTGG GCCTCCCACG AGAAGATGCA CGAGGGTGAC</p> <p>151 GATGCCCTG GCCACCACCA TAAGCCAGGC CTCGGGGAGG GCACCCCCTA</p> <p>201 AGACCACAGT GGACAAGATC ACAGTGGCCA CGGACACGGC CACAGTCATG</p> <p>251 GTGGCCACGG CCACAGCCAC TAATCAGGAG GCCAGGCCAC CCTGCCTCTA</p> <p>301 CCCAACCAAGG GCCCCGGGGC CTGTTATGTC AAACTGTCTT GGCTGTGGGG (<u>SEQ ID NO: 16</u>)</p>
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866	<p>1 NGATCTTCT AGGAGGGAGA CACTGGCCNC TCAAATCGTC CAGCGACCTT</p> <p>51 CCTCATCCAC CCCATCCCTC CCCAGTTCAT TGCACTTGA TTAGCAGCGG</p> <p>101 ACAAAGGAGT CAGACATT AAGATGGTGG CAGTAGAGGC TATGGACAGG</p> <p>151 GCATGCCACG TGGGCTCATA TGGGGCTGGG AGTAGTTGTC TTTCCTGGCA</p> <p>201 CTAACGTTGA GCCCCTGGAG GCACTGAAGT GCTTAGTGT CTTGGAGTAT</p> <p>251 TGGGGTCTGA CCCCAAACAC CTTCCAGCTC CTGTAACATA CTGGCCTGGA</p> <p>301 CTGTTTCTC TCGGCTCCCC ATGTGTCCTG GTTCCCGTT CTCCACCTAG</p> <p>351 ACTGTGAACC TCTCGAGGGC AGGGACCACA CCCTGTACTG TTCTGTGTCT</p> <p>401 TTCACAGCTC CTCCCACAAT GCTGAATATA CAGCAGGTGC TCAATAATG</p> <p>451 ATTCT (<u>SEQ ID NO: 17</u>)</p>
871	1 GCAAGTGTGT TGTGTTACAG TGTCACAACA CCGAG (<u>SEQ ID NO: 18</u>)
872	<p>1 GATCTCTCCC TACGCAAAAC GTATTGTAGT GAAAGGGTCT TCTTTACTAC</p> <p>51 CTTAATAAAA CAGCTAGTGT G (<u>SEQ ID NO: 19</u>)</p>

874	1 GATCTAAATA CAAAGGATAT ACAGTCTTGA ATCTAAAATA ATTTGCTAAC 51 TATTGTGATT CTTCAGAGAG AACTACTA (<u>SEQ ID NO: 20</u>)
876	1 GATCTAGTCC GGACATGCTG TGTATATTGT AACGTTAAAT GAAAAAAAGAA 51 CCCCCCTTG TATTATAGTC ATGCGGTCTT ATGTATGATA AACAGTTG (<u>SEQ ID NO: 21</u>)
878	1 GATCTTTGT AGTCACCTCT GTATCTTATG TCTGGTTGAG GGGTGCTTT 51 ACTTGTCTGG CATTGCATT CAATGATCTT TCAGTCATGT CAGTTAGACT 101 AAAAATTATT TCTG (<u>SEQ ID NO: 22</u>)
880	1 CCCAAGCCCC TTGGACACTG CAGCTCTTT CAGTTTTGC TTACACACAA 51 TTCATTCTTT GCAGCTAATT AAGCCGAAGA AGCGTGGGAA TCAAGTTGG 101 AACAGAGATT AAAAAGTTC TT (<u>SEQ ID NO: 23</u>)
881	1 GCTCTGGAGG ACAATCCAGG AACTACATTA CCTGGACTGT ATGCTGGTCA 51 TTTCTACAGA CAGCATTCAAG TATTGAGTG TACGGTAACT GTCTGGGTG 101 ATTCCCTATAA GATCATTATA CTG (<u>SEQ ID NO: 24</u>)

882	1 GATCTTCTC CTTGAATATC TTTCGATAAAA CAACAAAGGTG GTGTGATCTT 51 AATATATTG AAAAAAAACTT CATTCTCGTG AGTCATTAA ATGTGTACAA 101 TGTACACACT GGTACTTAGA GTTCTGTTT GATTCTTTT TAATAAACTA 151 C (<u>SEQ ID NO: 25</u>)
883	1 TGTCACTCAT GCCCTGGGAC TGCTTCTCCA GCCAGGCGGG CGCCATACGT 51 CCCACACTAG TGAAGGTCAA TGTCTCAGAA CAACACCTCT AT (<u>SEQ ID</u> <u>NO: 26</u>)
884	1 GATCTGGCCT GTTCCTGCGT CTGCGGAGCA GGCTTGTCT CCCAGCTATC 51 TATAACCTTA CCTAGAGTGT CGACTTGTGG GTTCCTGTTG CTGAGACTTC 101 CTGGATGGAG CCGCCCTCAC CGCCGGACCC GTAGCACTGC GCGGAAGTGT 151 GTCCAATAAA GT (<u>SEQ ID NO: 27</u>)
885	1 GATCTGATT GCTAGTTCTT CCTTGTAGAG TTATAAATGG AAAGATTACA 51 CTATCTGATT AATAGTTCT TCATACTCTG CATATAATT GTGGCTGCAG 101 AATATTGTAA TTTGTTGCAC ACTATGTAAC AAAACAACTG AAGATATGTT 151 TAATAAATAT TGTACT (<u>SEQ ID NO: 28</u>)

894	1 GATCTTATG AGAGCAGTAT TTTCTGTGTT TTCTTTAA TTTACAGCCT 51 TTCTTATTG GATATTTT TAATGTTGTG GATGAATGCC AGCTTCAGA 101 CAGAGCCCAC TTAGCTTGTC CACATGGATC TCAATGCCAA TCCTCCATTG 151 TTCCTCTCCA GATATTTTG GGAGTGACAA ACATTCTCTC ATCCTACTTA 201 GCCTACCTAG ATTTCTCATG ACGAGTTAAT GCATGTCCGT GGTGGGTGC 251 ACCTGTAGTT CTGTTATTG GTCA <u>(SEQ ID NO: 29)</u>
895	1 GATCTAAGTT AGTCCAAAAG CTAAATGATT TAAAGTCAAG TTGTAATGCT 51 AGGCATAAGC ACTCTATAAT ACATTAAATT ATAGGCCGAG CAATTAGGGA 101 ATGTTCTGA AACATTAAAC TTGTATTAT GTCACTAAAA TTCTAACACA 151 AACTAAAAAA ATGTGTCTCA TACATATGCT GTACTAGGCT TCATCATGCA 201 TTTCTAAATT TGTGTATGAT TTGAATATAT GAAAGAATT ATACACGAGT 251 GTTATTAAAA ATTATTAAAA ATAAATGTA <u>(SEQ ID NO: 30)</u>

896	1 GATCTTATAAG GCCTGTCTCA TCAGGGTGGT GTCAGCCCAG CTAGGATTAG 51 GCAGAATTGG GTGGGGGCTG TAGTGCACTT TTGGCACAGC ATGTACCTGT 101 CTGACTAATT CTCTGTCTT TCTTCCTGT TGCAATTCA TAGGCTTAGC 151 ATCTTCTGAA TGGTGTAG TAGGTCACTCC TGTTGATTTC CTGCTAGGGA 201 GTAGCATACT CTGGCTCTGT ACCACTGGCC AAGGGACTTA AGGATAGATG 251 AAGGGCTGCA GTTTGTTAA ATGGAACAAT ATGAAGAGA (<u>SEQ ID</u> <u>NO: 31</u>)
T10 3	1 GATCTTCTC CTTGAGTATC TTTGATAAAA CAACAAAGTG GTGTGATCTT 51 AATATATTTG AAAAAAAACTT CATTCTCGTG AGTCATTAA ATGTGTACAA 101 TGTACACACT GGTACTTAGA GTTTCTGTTT GATTCTTTT TAATAAACTA 151 C (<u>SEQ ID NO: 32</u>)
T10 4	1 GATCTCTGCT CATAGAATGC ATGGGGAGCC TTCCAGCTCA CTCTCCCTGA 51 GGACTGGCTT GACAGGGGCT ATGGGTTGC TTTGG (<u>SEQ ID NO: 33</u>)

T10 5	1 GATCTGCGCT TCCAGAGCGC AGCTATCGGT GCTTGCGAGG AGGCAAGTGA 51 GGCCTATCTG GTTGGCCTTT TTGAAGACAC CAACCTGTGT GCTATCCATG 101 CCAAACGTGT ACAATTATG CAAAAGACA TCCAGCTAGC ACGCCGCATA 151 CGTGGAGAAC GTGCTTAAGA ATCCACTATG ATGGGAAACA (<u>SEQ ID</u> <u>NO: 34</u>)
T10 7	1 GATCTAAATG TGAACAGTTT ACTAATGCAC TACTGAAGTT TAAATCTGTG 51 GCACAATCAA TGTAAGCATG GGGTTGTTT CTCTAAATTG ATTTGTAATC 101 TGAAATTACT GAACAACTCC TATTCCCATT TTTGCTAAAC TCAATTCTG 151 GTTTGGTAT ATATCCATT CAGCTTAATG CCTCTAATT TAATGCCAAC 201 AAAATTGGTT GTAATCAAAT TTTAAAATAA TAATAATTG GC (<u>SEQ ID</u> <u>NO: 35</u>)
T76	1 GCCTTTCGA TAGTTCGGG TCAGGTAAAA ATGGCCTCCT GGCGTAAGCT 51 TTTCAAGGTT TTTGGAGGC TTTTGTAAGG TTGTGATAGG AACTTGGAC 101 CTTGAACCTTA CGTATCATGT GGAGAAGAGC CAATTAAACA AACTAGGAAG 151 ATGAAAAGGG AAATTGTGGC CAAAACTTG GGAAAAGGAG GTTCTAAAA 201 TCAGTGTTC CCCTTT (<u>SEQ ID NO: 36</u>)

T8	<p>1 GATCTATGCA CAAGAACCCC TTTACCCAT GACCAACATC GCAGACACAT</p> <p>51 GTGCTGGCCA CCTGCTGAGC CCCAAGTGG ACGAGACAAG CAGCCCTTAG</p> <p>101 CCCTTCCCCT CTGCAGCTTC CAGGCTGGCG TGCAGCATCA GCATCCCTAG</p> <p>151 AAAGCCATGT GCAGCCACCA GTCCATTGGG CAGGCAGATG TTCCTAATAA</p> <p>201 AGCT (<u>SEQ ID NO: 37</u>)</p>
T81	<p>1 GATCTTCCT CCTGGTTACT GTGAAGCCTG TTGGTTGCT GCTGTCGTT</p> <p>51 TTGAGGAGGG CCCATGGGG TAGGAGCAGT TGAACCTGGG AACAAACCTC</p> <p>101 ACTTGAGCTG TGCCTAGACA ATGTGAATTCTGTGTTGCT AACAGAAGTG</p> <p>151 GCCTGTAAGC TCCTGTGCTC CGGAGGGAAG CATTCTGG TAGGCTTGA</p> <p>201 TTTTCTGTG TGTTAAAGAA ATTCAATCTA CTCATGATGT GTTATGCATA</p> <p>251 AACATTTCT GGAACATGGA TTTGTGTTCA CCTTAAATGT GAAAATAAT</p> <p>301 CCTA (<u>SEQ ID NO: 38</u>)</p>

T82	<p>1 ATCTTCCTC CTGGTTACTG TGAAGCCTGT TGGTTGCTG CTGTCGTTT</p> <p>51 TGAGGAGGGC CCATGGGGGT AGGAGCAGTT GAACCTGGGA ACAAACCTCA</p> <p>101 CTTGAGCTGT GCCTAGACAA TGTGAATTCC TGTGTTGCTA ACAGAAGTGG</p> <p>151 CCTGTAAGCT CCTGTGCTCC GGAGGGAAGC ATTCCTGGT AGGCTTGAT</p> <p>201 TTTCTGTGT GTAAAGAAA TTCAATCTAC TCATGATGTG TTATGCATAA</p> <p>251 AACATTCTG AACATGGAT TTGTGTCAC CTTAAATGTG AAAATAAAC</p> <p>301 CTATTTCTA TG (<u>SEQ ID NO: 39</u>)</p>
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T85	<p>1 GATCTTGGC AGGCCATTG GACTCTTGG GGTCACTGTC GCAATTCTTC</p> <p>51 ATACCTCCAG AGTGAAGATG GGTGACTAGA TGATATGTGT</p> <p>GGGTGGGCC</p> <p>101 GTGCCTCACT TTTATTATT GCTGGTTTC CTGGGACAGC TGGAGCTGTG</p> <p>151 TCCCTTAACC TTTCAGAGGC TTGGTGTCA GGGCCCTCCC TGCACCTCCCC</p> <p>201 TCTTGCTGCG TGTTGATTG GAGGCACTGC AGTCCAGGCC</p> <p>GAGTCCTCAG</p> <p>251 TGCAGGGAGC AGGCTGCTGC TGCTGACTCT GTGCAGCTGC</p> <p>GCACCTGTGT</p> <p>301 CCCCCACCTC CACCCCTCAAC CCATCTCCT AGTGTGTG AAATAAACTT</p> <p>351 GGTAT (<u>SEQ ID NO: 40</u>)</p>
T98	<p>1 GATCTTCCAC GTCTCCATCT CAGTACACAA TCATTTAATA TTTCCCTGTC</p> <p>51 TTACCCCTAT TCAAGCAACT AGAGGCCAGA AAATGGGCAA</p> <p>ATTATCACTA</p> <p>101 ACAGGTCTT GACTCAGGTT CCAGTAGTTC ATTCTAATGC CTAGATTCTT</p> <p>151 TTGTGGTTGT TGCTGGCCA ATGAGTCCCT AGTCACATCC</p> <p>CCTGCCAGAG</p> <p>201 GGAGTTCTTC TTTGTGAGA GACACTGTAA ACGACACAAAG</p> <p>AGAACAAAGAA</p> <p>251 TAAAAA (<u>SEQ ID NO: 41</u>)</p>

933

1 TTATATATT TTCTTAAATA TGTTTATTG TCTTCTCTAA GCAAAAAGTT
51 CTTAATAAAC ATAGTATTC TCTCTGCGTC CTATTCATT AGTGAAGACA
101 TAGTCACCT AAAATGGCAT CCTGCTCTGA ATCTAGACTT TTTAGAAATG
151 GCATATGTT TTGATGATAT GTCAACATTG AAAATAGTCC TAATTAAATT
201 GTTGGTTAAA TGTAATGTCA ACTCTTATA AACTTAAATA
TAAACAAAGTA
251 ATTAACCACT CTAAGTAATA AAACACATT CACCTGTGTT CTGAGTGTA
(SEQ ID NO: 42)

967

1 ATGAATCCTT GCCACCTCCA CCTGCAGAAC TGTTATAAAAT ATTACAACCTT

51 GCTTTTAGC TGATCTTCCA TCCTCAAATG ACTCTTTTT CTTTATATGT

101 TAACATATAT AAAATGGCAA CTGATAGTCA ATTTGATT
TTATTCAGGA

151 ACTATCTGAA ATCTGCTCAG AGCCTATGTG CATAGATGAA ACTTTTTTT

201 AAAAAAAAGTT ATTTAACAGT AATCTATTAA CTAATTATAG TACCTATCTT

251 TAAAGTATAG TACATTTAC ATATGTAAAT GGTATGTTTC AATAATTAA

301 GAACTCTGAA ACAATCTACA TATACTTATT ACCCAGTACA GTTTTTTTC

351 CCCTGAAAAG CTGTGTATAA AATTATGGTG AATAAACTTT
TATGTTCCA

401 TTCAAAGAC CAGGGTGGAG AGGAATAAGA GACTAAGTAT
ATGCTTCAAG

451 TTTAAATTA ATACCTCAGG TATTAAAATA AATATTCCAA
GTTTGTGGGA

501 AATGGGGAGA TTAAAATG (SEQ ID NO: 43)

978

1 TTATGTGGCC TTAGGTAGCT GGTTGTACAT CTTTCCCTAA ATCGATCCAT
51 GTTACCACAT AGTAGTTTA GTTTAGGATT CAGTAACAGT GAAGTGTAA
101 CTATGTGCAA CGGTATTGAA GTTCTTATGA CCACAGATCA
TCAGTACTGT
151 TGTCTCATGT AATGCTAAAA CTGAAATGGT CCGTGTTGC
ATTGTTAAAA
201 ATGATGTGTG AAATAGAATG AGTGCTATGG TGTTGAAAAC
TGCAGTGTCC
251 GTTATGAGTG CCAAAAATCT GTCTGAAGG CAGCTACACT
TTGAAGTGGT
301 CTTTGAATAC TTTAATAAAA TTTATTTGA TA (SEQ ID NO: 44)

981

1 TAGGTGAACC CTTATTCTGC AGGGTTCTCC CTCCCACCTT AAAGAAGTTC
51 CCCTTATGTG GGTTGCCTGG TGAATGGCCT TCCTTCCCGC CAGAGGGCTT
101 GTGAACAGAC CGGAGAGGAC AGTGGATTGT TTATACTCCA
GTGTACATAG
151 TGTAATGTAG CGTGTTACA TGTGTAGCCT ATGTTGTGGT CCATCAGCCC
201 CTCACATTCC TAGGGGTTTG AGATGCTGTA CGTGGTATGT
GACACCAAAG
251 CCACCTCTGT CATTGTTGT GATGTCTTT CTTGGCAAAA GCCTTGTGTA
301 TATTGTATA TTACACATT GTACAGAATT TTGGAAGATT TTCAGTCTAG
351 TTGCCAAATC TGGCTCCTTT ACAAAAG (SEQ ID NO: 45)

982	1 AGAATCTCTT ATGTTCTCAG AGGAAGGTGG AAGAAACCAT GGGCAGGAGT 51 AGGAATTGAG TGATAAACAA TTGGGCTAAT GAAGAAAACT TCTCTTATTG 101 TTCAGTCAT CCAGATTATA ACTTCAATGG GACACTTAG ACCATTAGAC 151 AATTGACACT GGATTAAACA AATTCACATA ATGCCAAATA CACAATGTAT 201 TTATAGCAAC GTATAATTG CAAAGATGGA CTTTAAAAGA TGCTGTGTAA 251 CTAAACTGAA ATAATTCAAT TACTTATTAT TTAGAATGTT AAAGCTTATG 301 ATAGTCTTT CTAATTCTTA ACACTCATAC TTGAAATCTT TCTGAGTTTC 351 CCCAGAAGAG AATATGGGAT TTTTTTGAC ATTTTGACT CATTAAATAA 401 TGCTCTTGTG TTTACCTAGT ATATGTAGAC TTTGTCTTAT GTGTCAAAAG 451 TCCTAGGAAA GTGGTTGATG TTTCTTATAG CAATTAAAAA TTATT <u>(SEQ</u> <u>ID NO: 46)</u>
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905	1 ATCTCAGTGA GCTGAGATCA CACCACTGCA CTCCAAGTGG GCGACAGAGC 51 AAGA <u>(SEQ ID NO: 47)</u>
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910	1 GATCTGTAAT TCAGGTGTTT TCTGTACAGC CATACTAGA TAATGAAGCC 51 AAAAGGCTTT TAATTACACC ATGGCCTAAA ATAAATTCA <u>CA (SEQ ID NO: 48)</u>
915	1 TATTTTCAG CTGAGTTATT AGGGAGTCAT TATTCTGTGG TACAATGCTG 51 CAAAAAGCAT CATGTGGAAG AATGGGAAC ATGCCTACAT TATGAAGTGA 101 TGTATAACAC AATGCAAATC TG <u>(SEQ ID NO: 49)</u>
916	1 GATCTTTTT CATAAAAAAA TGTTCAATT A TCAGGCCGGG TGCAGTGGGG 51 CTCATGCCTG TAATCCCAAC ACTTTGGGAG GCCGATGCAG GCGGATCACT 101 AGGTCAGCAG ATCGAGACCA TCCTGGCTAA CACAGTGAAA CCT <u>(SEQ ID NO: 50)</u>
921	1 GATCTTATT TTTAGCCATG CACTGTTGTG AGGAAAATT CCTGTCTTGA 51 CTGCCATGTG TTCATCATCT TAAGTATTGT AAGCTGCTAT GTATGGATT 101 AAACCGTAAT CATATCTTT TCCTATCTAT CTGAGGCACT GGTGGAATAA 151 AGAACCTGTA TATTTACTT TGTTGCAGAT AGTCTGCCG CATCTGGCA 201 AGTTGCAGAG A <u>(SEQ ID NO: 51)</u>

927	1 GATCTTCGTG AAGACCTGAC TGGTAAGACC ATCACCCCTCG AGGTGGAGCC 51 CAGTGACACC ATCGAGAATG TCAAGGCAAA GATCCAAGAT AAGGAAGGCA 101 TCCCTCCTGA TCAGCAGAGG TTGATCTTG CTGGGAAACA GCTGGAAGAT 151 GGACGCACCC TGTCTGACTA CAACATCCAG AAAGAGTCCA CTCTGCACTT 201 GGTCCCTGCGC TTGAGGGGGG GTGTCTAAGT TTCCCCTTT AAGGTTCAA 251 CAAATTTCAT TGCACCTTCC TTTCAATAAA GTTG <u>(SEQ ID NO: 52)</u>
928	1 GATCTTCCT CCTGGTTACT GTGAAGCCTG TTGGTTGCT GCTGTCGTTT 51 TTGAGGAGGG CCCATGGGG TAGGAGCAGT TGAACCTGGG AACAAACCTC 101 ACTTGAGCTG TGCCTAGACA ATGTGAATTCTGTGTTGCT AACAGAAGTG 151 GCCTGTAAGC TCCTGTGCTC CGGAGGGAAAG CATTCCCTGG TAGGCTTGA 201 TTTTCTGTG TGTTAAAGAA ATTCAATCTA CTCATGATGT GTTATGCATA 251 AAACATTCT GGAACATGGA TTTGTGTTCA CCTTAAATGT GAAAATAAAT <u>(SEQ ID NO: 53)</u>

930	<p>1 GATCTTCGG GTTCTCTCT CTAACTCAGC TCTTCGTCC CAGAAACCCA</p> <p>51 GATGTAATCC CCCTACGTGG TGCTTGGGGC ATCCCGATAC CATCTCAGTA</p> <p>101 AATCTCCTAC ATTGGCCTCC TCACCCTCCC CGGGACCCAC ACCCTTCAGG</p> <p>151 TCCTCACCCCT GAGACAGGAG GGACCCTCTG AGATCAGGGA CCCTTAGGTC</p> <p>201 TCACTGCTCT CTGATTATA GCTCAACTGG GCCCCCAGTT CCATACCCCA</p> <p>251 GCATTCCCGG TCACTCCCTC CCTAATCTGA GCATCACTCA AGCTCTTAT</p> <p>301 TAAACTC (<u>SEQ ID NO: 54</u>)</p>
939	<p>1 ATCTCTCTCC CTACGCAAAA CCCTATTGTA GTAAAAAAAGT CTTCTTTACT</p> <p>51 ATCTTAATAA AACAGATATT GTG (<u>SEQ ID NO: 55</u>)</p>
945	<p>1 ATCTATTCTT GTAGATTIT TTTGTGTGGG TCTATGTTTC ATTCACTCTGC</p> <p>51 TTTCAGGCTG GATTATAAC AAGCAGAACT TTTAAAACG (<u>SEQ ID NO: 56</u>)</p>
949	<p>1 GATCTAAATA TTTTCAGCT GAGTTATTAC GGAGTCATTA TTCTGTGGTA</p> <p>51 CAATGCTGCA AAAAGCATCA TGTGGAAGAA TGGGAACTAT GCTTACTTTA</p> <p>101 TGAAGTGATG TATAACACAA TGAAA (<u>SEQ ID NO: 57</u>)</p>

952	1 CTACCCCGTG ACTCAGTTAC CTCCCCTGG GTCCCTCCA CATCATGTGG 51 GAATTGTTAGG AGCTACAATT CAAGATGAGA TTTGGATGGG GTCACAGCCA 101 AACCATATCA CTGAGGTATC AAGGAGATTCTT <u>(SEQ ID NO: 58)</u>
954	1 GATCTGATT GCTAGTTCTT CCTTGTAGAG TTATAAATGG AAAGATTACA 51 CTATCTGATT AATAGTTCT TCATACTCTG CATATAATTGTGTT GTGGCTGCAG 101 AATATTGTAA TTTGTTGCAC ACTATGTAAC AAAACAACTG AAGATATGTT 151 TAATAAATAT TGTACTTATTG <u>(SEQ ID NO: 59)</u>
975	1 NGATCTTCT CCTTGAATAT CTTTCGATAA ACAACAAGGT GGTGTGATCT 51 TAATATATTG GAAAAAAACT TCATTCTCGT GAGTCATTAA AATGTGTACA 101 ATGTACACAC TGGTACTTAG AGTTCTGTT TGATTCTTT TTAATAAAA <u>(SEQ ID NO: 60)</u>

976	1 GATCTGCTAG AAGATGGTTT TGGAGAGCAC CCCTTTACC ACTGCCTGGT 51 TGCAGAAGTG CCGAAAGAGC ACTGGACTCC GGAAGGTAAC CCCTCGCCCT 101 TTCCAGAAC CAGAGAGACC AAGTGTATG TAAGAAGTAG TGTCGGCTGT 151 GTAGAACCA TGACTACACA GGCGAAGTT ACTGAGAACT TGGACAGAAA 201 AAATAGCCAG CAAGTGT <u>(SEQ ID NO: 61)</u>
984	1 CATTCACACA TTTAACCTCC TTCCATACCA AATCTT <u>(SEQ ID NO: 62)</u>
986	1 GATCTGGACA GCAGAATGTT ATAACGCAAG TTCATGTGTT GCTCCCAACT 51 CCATTCTCTT TTCTCTCGTG CAACCAGTTT GCCCATTCTC TTCCTATTAC 101 TTGCTC <u>(SEQ ID NO: 63)</u>
T11 3	1 TCAGAGATTT GCAAAGACTC ACGTTTTGT TGTTTCTCA TCATTCCATT 51 GTGATACTAA GAAACTAAGA AGCTTAATGA AAAGAAATAA AATGCCTATG <u>(SEQ ID NO: 64)</u>

T11	1 GATCTGCGCT TCCAGAGCGC AGCTATCGGT GCTTGCGAGG 6 AGGCAAGTGA
	51 GGCCTATCTG GTTGGCCTTT TTGAAGACAC CAACCTGTGT GCTATCCATG
	101 CCAAACGTGT ACAATTATG CAAAAGACA TCCAGCTAGC ACGCCGCATA
	151 CGTGGAGAAC GTGCTTAAGA ATCCACTATG ATGGGAAACA <u>(SEQ ID</u> <u>NO: 65)</u>
T12	1 GATCTGTGAA ATGCTATCTC TCCTGAAGCA ATACTGTTGA CCAGAAAGGA 3
	51 CACTCCATAT TGTGAAACCG GCCTAATTT TCTGACTGAT ATGGAAACGA
	101 TTGCCAACAC ATACTCTAC TTTAAATAA ACAACTTGA TGATGTAAC
	151 TGACCTCCA GAGTTATGGA AATTTGTCC CCATGTAATG AATAAATTGT
	201 ATGTAT <u>(SEQ ID NO: 66)</u>